

Please indicate:

MEDICARE FORM

Entyvio[®] (vedolizumab) Injectable Medication Precertification Request

Page 1 of 3

Start of treatment: Start date / /

(All fields must be completed and legible for precertification review.)

For Michigan MMP:FAX:1-844-241-2495PHONE:1-855-676-5772

For other lines of business: Please use other form.

Note: Entyvio is preferred on MA and MAPD plans.

Continuation of therapy	v: Date of last treatment/	/						
Precertification Requested By:		Phone:		Fax:				
A. PATIENT INFORMATION								
First Name:	L	.ast Name:						
Address:	C	City:		State:	ZIP:			
Home Phone:	Work Phone:		Cell Phone:					
DOB: Allergies:			Email:					
Current Weight: lbs or	kgs Height	t: ir	ches or		cms			
B. INSURANCE INFORMATION								
Aetna Member ID #:	Does patient have of	Does patient have other coverage?						
Group #:	If yes, provide ID#: _	Ca	rrier Name: _					
Insured:	Insured:							
C. PRESCRIBER INFORMATION								
First Name:	Last Name:		(Check Or	ne): 🗌 M.D. 🗌] D.O. 🗌 N.P. 🗌 P.A			
Address:		City:		State:	ZIP:			
Phone: Fax:	St Lic #:	NPI #:	DEA #:	UI	PIN:			
Office Contact Name:			Phone:					
D. DISPENSING PROVIDER/ADMINISTRATION	INFORMATION							
ce of Administration: Self-administered Physician's Office Outpatient Infusion Center Phone: Center Name: Home Infusion Center Phone: Agency Name: Administration code(s) (CPT): dress:		_ ☐ Other: - Name: - Address: - City:	fice macy	└── Retail Pharmacy └── Mail Order				
City: State: _		Phone:		Fax:				
Phone: Fax:		- TIN:		PIN:				
NPI:		NPI:						
E. PRODUCT INFORMATION								
Request is for Entyvio (vedolizumab): Dose:	Freque	ency:		HCPCS Code	e:			
F. DIAGNOSIS INFORMATION – Please indicate	primary ICD Code and specify any	y other where applicable.						
Primary ICD Code: Secondary ICD Code:		Other ICD Code:						
G. CLINICAL INFORMATION – Required clinical	information must be completed in i	ts <u>entirety</u> for all pr <u>ecertif</u>	ication reques	ts.				
For Initiation Requests (clinical documentation	required):							
Note: Entyvio is preferred on MA and MAPD plans.								
□ Yes □ No Has the patient had prior therapy □ Yes □ No Will Entyvio (vedolizumab) be used		-	c DMARDs (e	.g., adalimumab	, infliximab)?			
E. PRODUCT INFORMATION Request is for Entyvio (vedolizumab): Dose: F. DIAGNOSIS INFORMATION – Please indicate Primary ICD Code: G. CLINICAL INFORMATION – Required clinical For Initiation Requests (clinical documentation Note: Entyvio is preferred on MA and MAPD pl YesNo Has the patient had prior therapy	Freque primary ICD Code and specify any Secondary ICD Code: information must be completed in i required): lans. with Entyvio (vedolizumab) within f	NPI:	Other ICD Co	_ HCPCS Code ode: ts.	e:			

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB				
G. CLINICAL INFORMATION (continued) – R	equired clinical information must be comple	eted in its <u>entirety</u> for all precert	ification requests.				
Crohn's Disease							
	nosis of fistulizing Crohn's disease? <i>If yes,</i>		diagnosis: / /				
→ Please indicate the severity of the patient's Crohn's disease: ☐ Mild ☐ Moderate ☐ Severe ☐ Yes ☐ No Is there clinical evidence that the disease is active?							
\top \top $-$	 Is the Crohn's disease manifested by at 	least one of the following?					
	\rightarrow Check all that apply: \Box abdominal pain		diarrhea 🔲 internal fistulae				
	intestinal obstruction megacolon						
☐ Yes ☐ No Was treatmer	nt with corticosteroids ineffective?						
	o Was treatment with corticosteroids not tol	erated or contraindicated?					
	\rightarrow \Box not tolerated \Box contraindicated						
		rticosteroids was tried? [_] hydi Please explain:	rocortisone				
Which of the	following corticosteroids was tried?	ocortisone	lone				
			ain:				
	nt with 6-mercaptopurine (6-MP) ineffective?						
$ \qquad \qquad$	o Was treatment with 6-mercaptopurine (6-	MP) not tolerated or contraindic	ated?				
	\rightarrow \Box not tolerated \Box contraindicated in with azathioprine ineffective?						
	o Was treatment with azathioprine not toler	rated or contraindicated?					
	\rightarrow \Box not tolerated \Box contraindicated						
Ulcerative Colitis							
Yes No Is the patient hospitalized fulm							
	the patient's ulcerative colitis: Mild nce that the disease is active?	Moderate 🗋 Severe					
	refractory to immunosuppression with cortic	costeroids (e.g., hydrocortisone	e, methylprednisolone, prednisone)?				
	o Does the patient require continuous imn						
	methylprednisolone, prednisone)?	2					
	Name and dose: Name: Please indicate the route: Oral I	Dose:					
Name and do		Dose:					
	te the route:						
	nt with immunosuppressant agent (e.g., aza						
	 Was treatment with immunosuppressan or contraindicated? 	t agent (e.g., azathioprine, m6-	mercaptopurine) not tolerated				
	\rightarrow \Box not tolerated \Box contraindicated						
	Provide the name of the drug	ug(s):					
	name of the drug(s):						
	nt with 5-aminosalicylic acid agents (e.g., ba						
	 Was treatment with 5-aminosalicylic acid not tolerated or contraindicated? 	a agents (e.g., baisalazide, me	salarinite, suitasalazitte)				
	\rightarrow \Box not tolerated \Box contraindicated						
	Provide the name of the dru	ug(s):					
Provide the n	ame of the drug(s): ne patient exhibit: more than 10 stools p	or day. 🗖 continuous blooding	a Dabdominal pain D distansion				
		mptoms, including fever and ar					
For Continuation requests (clinical document		inplome, melading lever and a					
	used concomitantly with aprelimast, tofacit	inib, or other biologic DMARDs	(e.g., adalimumab, infliximab)?				
	result of the patient receiving samples of Er	_					
Yes No Is there clinical documentation	n supporting disease stability?						
Yes No Is there clinical documentation	n supporting disease improvement?						
	vio (vedolizumab) within the past 6 months						
	tient have a documented severe and/or pote	entially life-threatening adverse	event that occurred during or				
•	previous infusion?	and through pro-modication in	the home or office astting?				
	No Could the adverse reaction be mana	yea unough pre-medication in	the nome of onice setting?				



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H. ACKNOWLEDGEMENT					
Request Completed By (Signature Required):			Date:	1	/

Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.

The plan may request additional information or clarification, if needed, to evaluate requests.